



ZETA RESEARCH

Tailoring research with you
Il tuo partner per la ricerca

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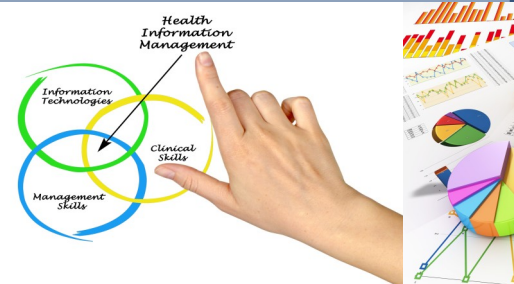


ZETA RESEARCH SRL

Zeta Research has qualified staff for the full management of clinical trials, at national and international level.


Zeta Research, active since 2002, is a **Contract Research Organization (CRO)**, certified at the Italian Drug Agency (AIFA), for the **statistical analysis** and **data management** of **pharmacological interventional studies**, according to DM 15 november 2011.

To offer a complete service to its customers, Zeta Research has set up 10 CATI stations for recall and phone services to patients and PI.



Today, Zeta Research has extensive experience in the design, development, and management of clinical trials, scientific studies and research projects, ensuring compliance with the international regulations, to obtain quality and independent results. The multidisciplinary approach of Zeta Research allows responding the specific customer needs, by using competent human resources.

The Company supports public and private clients such as multinational pharmaceutical companies, electro-medical and food enterprises, healthcare organizations, medical associations, European Authorities and partners in high level consortiums for the development of European Calls.



The services range from the creation of the **study design** with the drafting of a **protocol** and CRF to **document management**; from the selection of the Centers to the submission of the study to the **Ethics Committees** and notification to Competent Authorities; from the implementation of a dedicated site or portal for **data collection and management**; from remote or on-site **monitoring** to **data cleaning** or recoding (according to specific MedDRA - Medical Dictionary for Regulatory Activities); from the **statistical analysis** to the drafting of **scientific material** for publications.



In recent years, Zeta Research has focused its expertise and resources in the collection of online data (Electronic Data Capture - EDC) with the use of computer systems validated according to the Computer System Validation (CSV) in Clinical Data Management (CDM). The service guarantees results that can be used for registration or markings of new products, for the post-marketing surveillance and are expendable in the scientific community with publications or presentations.

Data obtained with the procedures developed by Zeta Research meet the specific regulatory requirements and international guidelines (application of ICH GCP, Annex 11 CFR part 11, Statement of Privacy, and standards SCDM, GCDMP & CDISC).



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